

**K170242 MultiGen™ 2 RF Generator System**May 25, 2017  
119 days to decisionK170242 · Product code: **GXD** · Neurology  
Source: <https://www.510kdatabase.net/k170242/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Generator, Lesion, Radiofrequency (GXD)
Date received	Jan 26, 2017
Decision date	May 25, 2017
Days to decision	119 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Stryker Corporation</b>
Location	Malwah, NJ, US
Contact	BECKY DITTY
Website	<a href="http://www.stryker.com/">http://www.stryker.com/</a>
510(k) history	81 submissions · 81 cleared · 2010-2023

Stryker Corporation is an American multinational medical technology company headquartered in Portage, Michigan. The company develops and markets surgical equipment, neurotechnology, orthopedic implants, and patient safety systems used globally across medical specialties. Stryker has received FDA 510(k) clearances from total submissions between 2010 and 2023. The company's cleared devices span orthopedic surgery, neurosurgery, general and plastic surgery, and ear, nose, and throat specialties. This regulatory record reflects the company's broad portfolio across surgical an...

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